What is claimed is:

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1. A method for producing an optically active  $\beta\text{-amino}$  acid of formula (2),

$$\begin{array}{c|c}
NH_2 \cdot bX' \\
0 \\
R^1 \times R^2
\end{array}$$
(2)

,wherein b is 0 or 1; the symbol \* shows that the carbon atom is a chiral carbon; R1 is a hydrogen atom, an alkyl group, a substituted alkyl group, a cycloalkyl group, a substituted cycloalkyl group, an aralkyl group, substituted aralkyl group, an aryl group, a substituted aryl group, an aliphatic heterocyclic group, a substituted aliphatic heterocyclic group, an aromatic heterocyclic group, a substituted aromatic heterocyclic group, an alkoxy group, a substituted alkoxy group, an aralkyloxy group, a substituted aralkyloxy group, an aryloxy group or a substituted aryloxy group; R<sup>2</sup> is a hydrogen atom, an alkyl group, a substituted alkyl group, a cycloalkyl group, a substituted cycloalkyl group, an aralkyl group, a substituted aralkyl group, an aryl group, a substituted aryl group, an aliphatic heterocyclic group, a substituted aliphatic heterocyclic group, an aromatic heterocyclic group, a substituted aromatic heterocyclic group, an alkoxy group, a substituted alkoxy group, an aralkyloxy group, a substituted aralkyloxy group, an aryloxy group, a substituted aryloxy group, an alkyloxycarbonyl group or an aralkyloxycarbonyl group; R3 is an alkoxy group, a

substituted alkoxy group, an aralkyloxy group, a substituted aralkyloxy group, an aryloxy group, a substituted aryloxy group, an amino group or a substituted amino group, X' is an acid, and  $R^1$  and  $R^2$  or  $R^2$  and  $R^3$  may be combined together to form a ring provided that  $R^1$  and  $R^2$  are not a hydrogen atom simultaneously, which comprises subjecting an enamine of formula (1),

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$$R^{1} \xrightarrow{R^{2}} R^{3}$$
 (1)

- , wherein  $R^1$ ,  $R^2$ ,  $R^3$  and X' have the same meanings as described above, and a is 0 or 1, to an asymmetric hydrogenation.
  - 2. The method as claimed in claim 1, wherein the asymmetric hydrogenation is carried out in the presence of an acid.
    - 3. The method as claimed in claim 1, wherein the asymmetric hydrogenation is carried out in the presence of a fluorine-containing aliphatic alcohol.
- 4. The method as claimed in any of claims 1 to 3,
  wherein the asymmetric hydrogenation in carried out in the presence of a catalyst for the asymmetric hydrogenation.
  - 5. The method as claimed in claim 4, wherein the catalyst for the asymmetric hydrogenation is a transition metal complex.
- 25 6. The method as claimed in claim 5, wherein the

transition metal complex is a complex of a metal which belong to the eighth group of the periodic table.

7. The method as claimed in either claim 5 or claim 6, wherein the transition metal complex has a chiral ligand.

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- 8. The method as claimed in claim 7, wherein the chiral ligand is a chiral phosphine ligand.
- 9. The method as claimed in claim 1, wherein the asymmetric hydrogenation is carried out in the presence of an acid and a fluorine-containing aliphatic alcohol.